



## Authenticity and Traceability of Vanilla Flavour by Analysis of Stable Isotopes

Hansen, Anne-Mette Sølvbjerg; Fromberg, Arvid; Frandsen, Henrik Lauritz

*Publication date:*  
2014

*Document Version*  
Publisher's PDF, also known as Version of record

[Link back to DTU Orbit](#)

*Citation (APA):*  
Hansen, A-M. S., Fromberg, A., & Frandsen, H. L. (2014). *Authenticity and Traceability of Vanilla Flavour by Analysis of Stable Isotopes*. Poster session presented at Food Integrity & Traceability Conference (ASSET 2014), Belfast, Ireland.

---

### General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.



# Authenticity and Traceability of Vanilla Flavour by Analysis of Stable Isotopes

Anne-Mette Sølvbjerg Hansen, Arvid Fromberg, Henrik Lauritz Frandsen

National Food Institute. Technical University of Denmark. Mørkhøj Bygade 19. DK-2860 Søborg. Denmark  
Corresponding author: asoha@food.dtu.dk

How can you be sure that the vanilla flavour added to your food product is actually a natural vanilla flavour originating from Madagascar as indicated on the label? In order to protect consumers against mislabeling and local producers against counterfeits, it is necessary to develop methods that can verify authenticity.

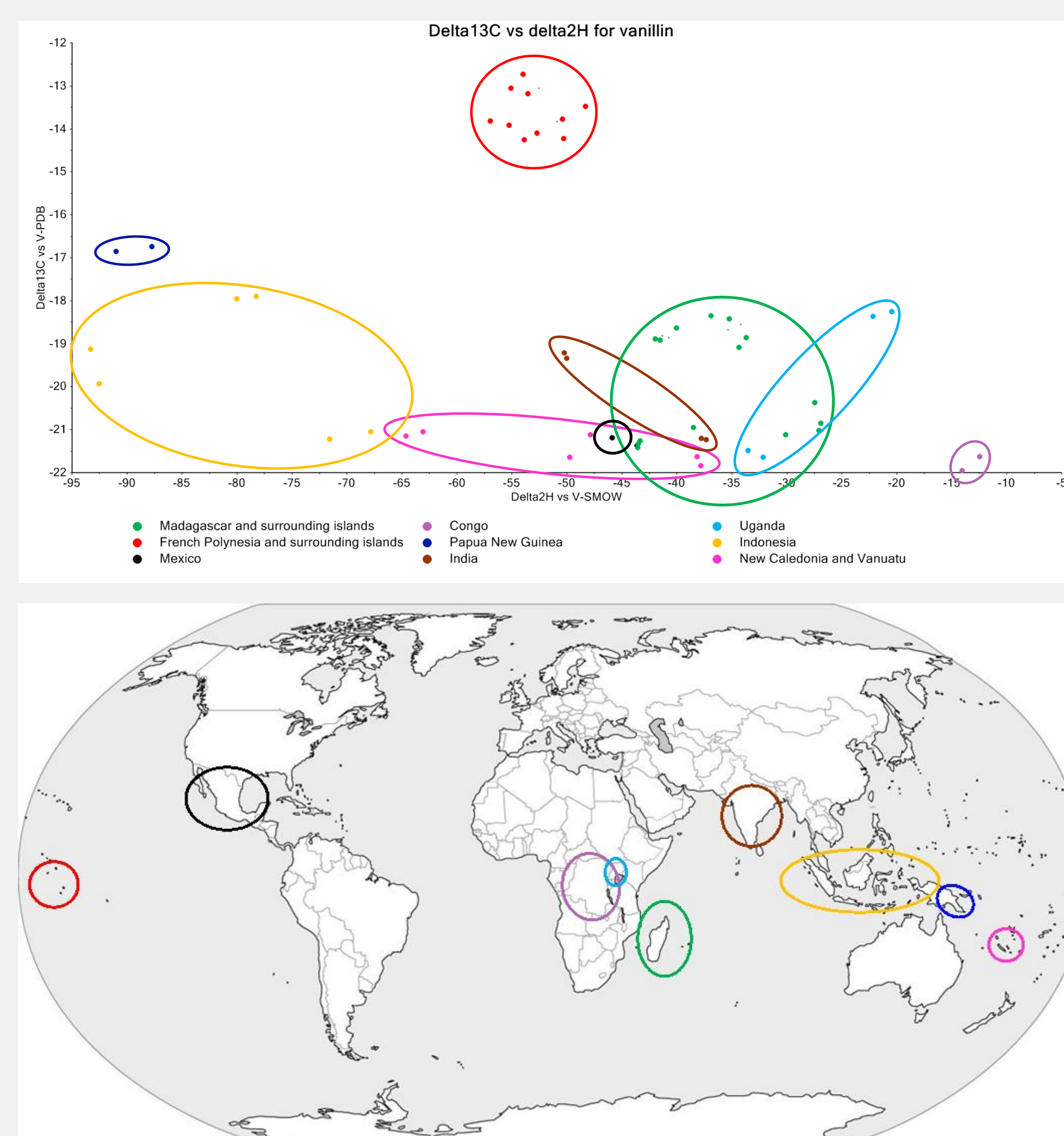
## Introduction

The characteristic flavour from vanilla pods is mainly due to 4-hydroxy-3-methoxybenzaldehyde, also called vanillin. Vanillin is added to a wide range of food products to obtain vanilla flavour. In general there are three types of vanillin: **natural vanillin** extracted from the vanilla orchid, **synthetic vanillin**, and **biovanillin** made by fermentation of e.g. eugenol. Because the vanilla plant uses the Crassulacean Acid Metabolism (CAM) for CO<sub>2</sub> fixation, the <sup>13</sup>C/<sup>12</sup>C ratio will differ from plants used as substrates for fermentations, as they use the Calvin Cycle (C3) for carbon fixation. Synthetic vanillin is made from petrochemicals which often are more depleted in the heavy carbon isotopes compared to CAM plants. Therefore the differences in ratios of stable carbon isotopes of vanillin were, in this study, used to determine the method of production.

Stable isotope ratios of hydrogen was used to trace the geographical origin of a vanilla plant. The ratio of <sup>2</sup>H/<sup>1</sup>H in precipitation vary due to longitudinal, altitudinal, and continental effects and this isotopic variation is incorporated into the plants' metabolites including vanillin.

## Results

$\delta^{13}\text{C}$  values of natural vanillin was significantly different ( $P < 0.001$ ) from the values of synthetic vanillin and biovanillin, while there was no difference between synthetic vanillin and biovanillin. Furthermore, a significant difference ( $P < 0.001$ ) of  $\delta^{13}\text{C}$  for the two types of natural vanilla pods was found: *Vanilla planifolia* and *Vanilla tahitensis*.



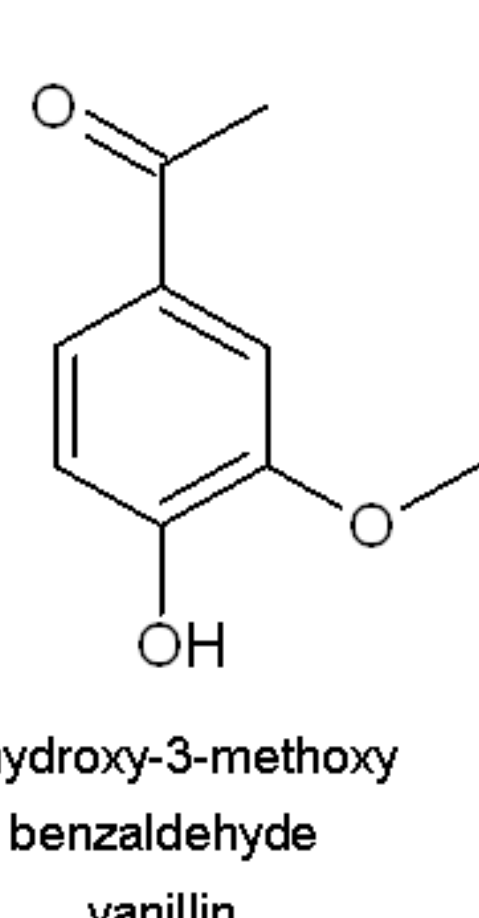
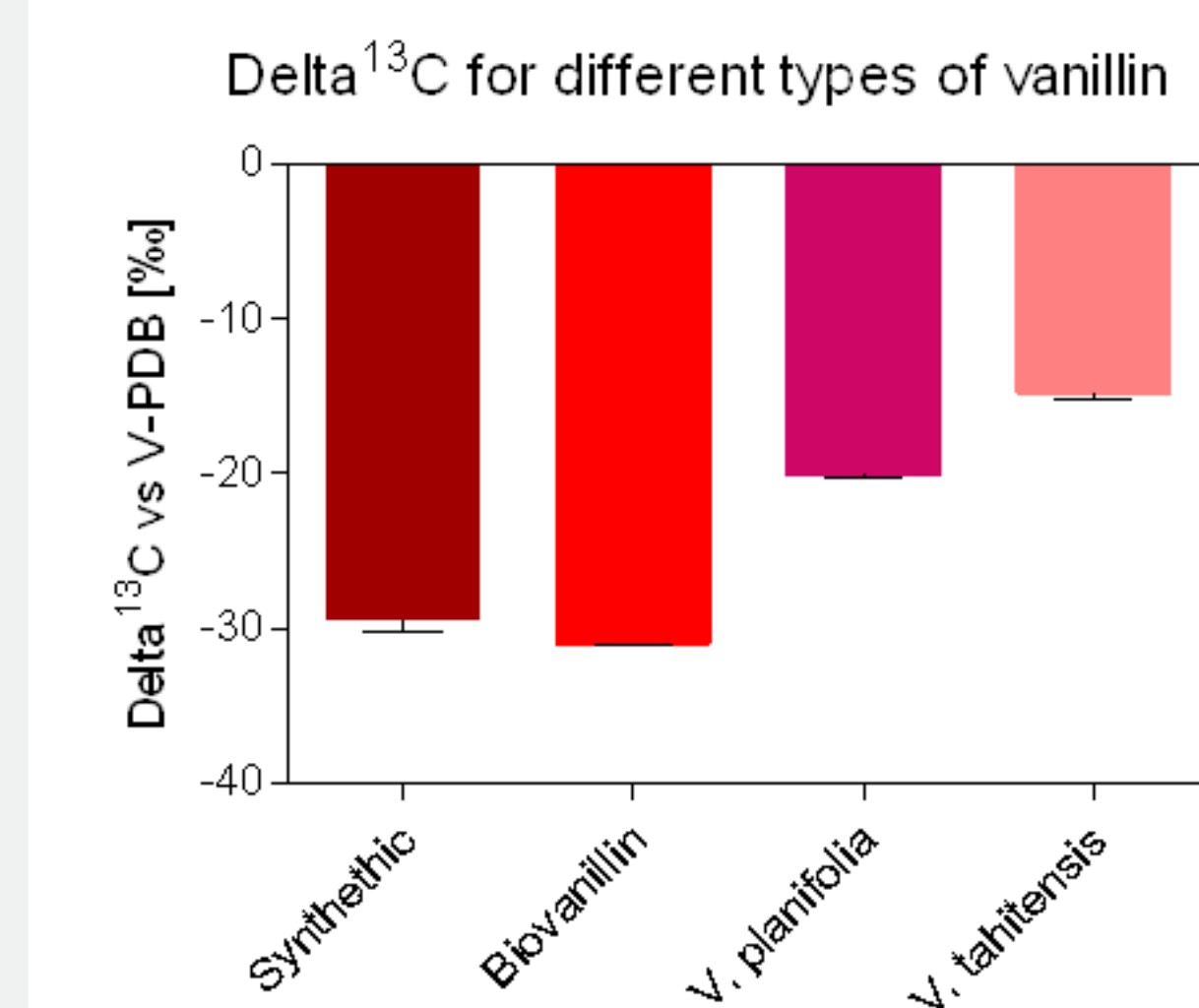
Stable isotopes of carbon and especially hydrogen can provide information about the geographical origin of vanilla pods. There is a tendency towards a depletion of <sup>2</sup>H in areas with high precipitation.

## Materials and methods

Vanillin was extracted from vanilla pods using EtOH/H<sub>2</sub>O (1:1) for 72h. After filtration the extract was further extracted with ethyl acetate/cyclohexane (1:1).

GC-IRMS: Trace GC Ultra fitted with a DB-5 capillary column (Agilent Technologies, Böblingen, Germany) (30m x 0.250mm i.d., d<sub>f</sub> 0.25 μm) coupled to Delta V Advantage Isotope Ratio Mass Spectrometer (Thermo Scientific, Bremen, Germany).

For measurements of  $\delta^{13}\text{C}$  a **Combustion Reactor** (NiO tube with NiO/CuO/Pt) was operated at 1000 °C. For determinations of  $\delta^2\text{H}$  a **High Temperature Conversion Reactor** consisting of a ceramic tube with no catalyst was operated at 1420 °C.



Values of  $\delta^{13}\text{C}$  for synthetic, bio and natural vanillin. Number of samples analyzed was 2 synthetic, 1 biovanillin, 28 *V. planifolia* and 10 *V. tahitensis*. Results are stated as  $\delta$  values:  $\delta^{13}\text{C} = ((R_{\text{sample}}/R_{\text{standard}}) - 1) \times 1000$ , where R is the ratio of the heavy to the light isotope.

